Application No. National Phase of PCT/IT/2006/000049 Amendment dated September 1, 2006 Preliminary Amendment Accompanying New Application

IN THE CLAIMS

1. (Original) Compound having the general formula (I):

or its geometric isomers, its optically active forms, diastereoisomers, its racemic forms, or its pharmaceutically acceptable salts, wherein R¹ is selected from the group consisting of: C₂-C₉ alkandiamine, C₂-C₆ amine; X is selected from the group consisting of: -S-S-, -S-, -CH₂-, -CH₂-CH₂-; m is an integer greater than zero and lower than eight; Ar represents an aromatic group; R¹ comprises a nitrogen linked directly to the carbonyl.

- 2. (Original) Compound according to claim 1, wherein X represents -S-S-.
- 3. (Currently amended) Compound according to claim 1 or 2, wherein m is an integer greater than two and lower than five.
 - 4. (Original) Compound according to claim 3, wherein m is four.

5. (Currently amended) Compound according to one of the previous claims claim 1, wherein Ar presents a formula selected from the group consisting of:

wherein R^5 is selected from the group consisting of: hydrogen, amine, nitroalkyl, -NH₂, nitro, halogen, hydroxy; R^6 is selected from the group consisting of: hydrogen, amine, alkandiamine, -NH₂; R^7 is selected from the group consisting of: hydrogen, group having an electron attractor inductive effect; R^{13} , R^{14} , R^{15} , R^8 and R^9 are selected, each independently of the others, from the group consisting of: hydrogen, hydroxy, halogen, alkoxy, alkyl, nitroalkyl, cyanoalkyl, nitro, cyano; R^{10} and R^{11} , are selected, each independently of the other, from the group consisting of: hydrogen, C_1 - C_4 alkyl; R^{12} represents a C_1 - C_4 alkyl; Y is selected from the group consisting of -CH- and -N-.

6. (Original) Compound according to claim 5, wherein Ar presents a formula selected from the group consisting of:

7. (Original) Compound according to claim 6, wherein Ar presents the formula:

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wherein R¹ represents a C₂-C₆ amine.

- 8. (Original) Compound according to claim 7, wherein R^1 presents the formula $-N(CH_2)_n$, wherein the nitrogen is directly linked to the carbonyl and n is an integer greater than one and smaller than five.
- 9. (Original) Compound according to claim 8, wherein n is three; R^{10} and R^{11} represent, each, a respective methyl; R^{12} represents an ethyl and is linked at the meta position with respect to the oxygen.
- 10. (Original) Compound according to claim 9, and having the following formula:

11. (Original) Compound according to claim 6, wherein Ar presents the formula:

wherein Y represents N, R¹ represents an alkandiamine having the formula –NR³-R²-NR⁴-; R² represents a C₂-C₅ alkyl; R³ and R⁴ are selected, each independently of the other, from the group consisting of: hydrogen, methyl; R¹³, R¹⁴, R¹⁵ are selected, each independently of the others, from the group consisting of: hydrogen, hydroxy, halogen, C₁-C₄ alkyl.

- 12. (Original) Compound according to claim 11, wherein R² represents a linear propyl; R³ and R⁴ each represent a hydrogen; R¹³ represents a halogen; R¹⁴ and R¹⁵ are selected, each independently of the other, from the group consisting of: halogen, hydroxy, C₁-C₄ alkoxy.
- 13. (Currently amended) Compound according to claim 11 or 12, wherein R¹³ represents a chlorine; R¹⁴ and R¹⁵ represent, each, a respective methoxy.
- 14. (Original) Compound according to claim 5, wherein Ar presents the formula:

 R^7 is selected from the group consisting of: hydrogen, C_1 - C_4 alkoxy, halogen; R^6 is selected from the group consisting of: $-NH_2$, alkandiamine, amine; R^1 represents a C_1 amine.

15. (Original) Compound according to claim 14, wherein R^6 is selected from the group consisting of: $-NH_2$ and amine C_1-C_4 .

16. (Original) Compound according to claim 14, wherein R⁷ is a chlorine situated in position 6; R⁶ represents –NH₂; R¹ represents –NH-CH₂-, wherein the nitrogen is linked to the carbonylic carbon.

17. (Original) Compound according to claim 5, wherein Ar presents the formula:

$$R^7$$
 R^5

wherein R¹ represents a C₂-C₆ alkandiamine.

- 18. (Original) Compound according to claim 17, wherein R^1 represents a C_3 - C_4 alkandiamine.
- 19. (Currently amended) Compound according to claim 17 or 18, wherein R^1 presents the formula $-NR^3-R^2-NR^4$, wherein R^2 represents a C_2-C_4 alkyl, R^3 and R^4 are selected, each independently of the other, from the group consisting of: hydrogen, methyl.
- 20. (Original) Compound according to claim 19, wherein R³ and R⁴ represent, each, a respective hydrogen.
- 21. (Currently amended) Compound according to claim 19 or 20, wherein R^2 represents $-(CH_2)_3$ -.
- 22. (Currently amended) Compound according to one of the claims from claim 17 to 21, wherein R⁷ represents a group having an electron withdrawing inductive effect.
- 23. (Original) Compound according to claim 22, wherein R⁷ is selected from the group consisting of: halogen, C₁-C₄ alkoxy.
- 24. (Original) Compound according to claim 23, wherein R⁷ represents a halogen.

25. (Currently amended) Compound according to one of the claims from claim 17 to 21, wherein R⁷ is selected from the group consisting of: halogen, hydrogen, methoxy; R⁵ is selected from the group consisting of: hydrogen, amine, nitroalkyl, halogen, hydroxy.

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- 26. (Currently amended) Compound according to one of the claims from claim 17 to 25, wherein R⁷ is situated in position 6.
- 27. (Currently amended) Compound according to one of the claims from claim 17 a 26, wherein R⁵ is selected from the group consisting of: hydrogen, C₁-C₄ amine, C₁-C₄ nitroalkyl, -NH₂, nitro, halogen.
- 28. (Currently amended) Compound according to one of the claims from claim 17 to 26, wherein R⁵ is selected from the group consisting of: hydrogen, halogen.
- 29. (Original) Compound according to claim 28, and having the following formula:

30. (Original) Compound according to claim 29, in form R:

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31. (Original) Compound according to claim 6, wherein Ar presents the formula:

wherein R¹ represents a C₃-C₉ alkandiamine.

- 32. (Original) Compound according to claim 31, wherein R^1 represents a C_6 - C_8 alkandiamine.
- 33. (Currently amended) Compound according to claim 31 or 32, wherein R^1 presents the formula $-NR^{16}-R^{17}-NR^{18}-R^{19}$, wherein R^{19} is linked to Ar and $-NR^{16}$ is linked to the carbonylic carbon; R^{17} is a C_2 - C_7 alkyl; R^{16} and R^{18} are selected, each independently of the other, from the group consisting of: C_1 - C_3 alkyl, hydrogen; R^{19} represents a C_1 - C_3 alkyl.
- 34. (Original) Compound according to claim 33, wherein R¹⁷ is a C₃-C₆ alkyl; R¹⁶ represents a hydrogen; R¹⁸ is selected from the group consisting of: ethyl, methyl, hydrogen; R¹⁹ represents a methyl.
- 35. (Currently amended) Compound according to one of the claims from claim 31 a 34, wherein R⁹ is selected from the group consisting of: hydrogen, hydroxy, halogen, C₁-C₄ alkoxy; R⁸ is selected from the group: hydroxy, halogen, C₁-C₄ alkoxy.
- 36. (Original) Compound according to claim 35, wherein R⁹ represents a hydrogen and R⁸ represents a methoxy situated in ortho or meta position with respect to the remaining part of the compound.

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37. (Original) Compound according to claim 36, and having the following formula:

- 38. (Currently amended) Compound having the general formula (I), as defined in any one of the claims from claim 1 to 37, for use as a medicament.
 - 39. (Cancelled)
 - 40. (Cancelled)
 - 41. (Cancelled)
- 42. (Currently amended) Pharmaceutical preparation comprising a compound having general formula (I), as defined in any one of the claims from claim 1 to 37, or a pharmaceutically acceptable salt, and an excipient and/or pharmaceutically acceptable diluent.
- 43. (Currently amended) Method for the treatment of Alzheimer's disease in a mammal; the method comprises comprising administering to said mammal an efficacious quantity of a compound having general formula (I), as defined in any one of the claims from claim 1 to 37.

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44. (Currently amended) Method for the synthesis of a compound having general formula (I), as defined in any one of the claims from claim 1 to 37, comprising an addition phase wherein a compound having the general formula (II):

is reacted with a compound having the general formula (III):

in basic conditions.

45. (New) Method for the treatment of a pathology characterized by deposits of β -amiloid (A β) in mammals comprising administering to said mammal an efficacious quantity of a compound having a general formula (I), as defined in claim 1.